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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/589,276	03/10/2008	Erich Wanker	009848-0356193	5727
	7590 08/16/201 VINTHROP SHAW PI		EXAM	IINER
ATTENTION: DOCKETING DEPARTMENT			SAMALA, JAGADISHWAR RAO	
P.O BOX 10500 McLean, VA 22			ART UNIT	PAPER NUMBER
			1618	
			MAIL DATE	DELIVERY MODE
			08/16/2010	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)					
Office Action Comment	10/589,276	WANKER ET AL.					
Office Action Summary	Examiner	Art Unit					
	JAGADISHWAR R. SAMALA	1618					
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence ad	ldress				
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period w - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 16(a). In no event, however, may a reply be tim ill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONEI	J. lely filed the mailing date of this co ○ (35 U.S.C. § 133).					
Status							
1) Responsive to communication(s) filed on							
	- action is non-final.						
3) Since this application is in condition for allowan	ice except for formal matters, pro	secution as to the	e merits is				
closed in accordance with the practice under E							
Disposition of Claims							
4)⊠ Claim(s) <u>1-22</u> is/are pending in the application.							
· · · · · · · · · · · · · · · · · · ·	4a) Of the above claim(s) is/are withdrawn from consideration.						
5) Claim(s) is/are allowed.							
6) Claim(s) is/are rejected.							
7) Claim(s) is/are objected to.							
8) Claim(s) 1-22 are subject to restriction and/or e	election requirement.						
Application Papers							
9) The specification is objected to by the Examine	-						
10) The drawing(s) filed on is/are: a) acce		Examiner.					
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).							
Replacement drawing sheet(s) including the correcti			FR 1.121(d).				
11) The oath or declaration is objected to by the Ex							
Priority under 35 U.S.C. § 119							
12) ☐ Acknowledgment is made of a claim for foreign	priority under 35 LLS C & 110(a)	-(d) or (f)					
a) ☐ All b) ☐ Some * c) ☐ None of:	priority under 35 0.5.6. § 119(a)	-(a) or (i).					
1. ☐ Certified copies of the priority documents	s have been received						
2. ☐ Certified copies of the priority documents		on No					
	• •	<u></u>	Stane				
<u> </u>	3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).						
* See the attached detailed Office action for a list of		d					
		u.					
Attachment(s)							
1) Notice of References Cited (PTO-892)	4) Interview Summary	(PTO-413)					
2) Notice of Draftsperson's Patent Drawing Review (PTO-948)	Paper No(s)/Mail Da	ite					
3) Information Disclosure Statement(s) (PTO/SB/08)	5) Notice of Informal P	atent Application					
Paper No(s)/Mail Date	o) 🔲 Otilet						

DETAILED ACTION

Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 1-7 (in part) and claim 16, drawn to pharmaceutical or diagnostic composition comprising one or more active substances wherein the one or more active substance is/are selected from a group consisting of If this group is elected, a further election of a single species of active substance with every component named is required.

Group II, claim(s) 1-7 (in part)and claim 8, drawn to pharmaceutical or diagnostic composition according to claim 1, wherein the active substance with a structure. If this group is elected, a further election of a single species of active substance with every component named is required.

Group III, claim(s) 1-7 (in part) and claim 9, drawn to pharmaceutical or diagnostic composition according to claim 1, wherein the active substance with a structure to formula I-1 is selected from. If this group is elected, a further election of a single species of active substance with every component named is required.

Group IV, claim(s) 1-7 (in part) and claim 10, drawn to pharmaceutical or diagnostic composition according to claim 1, wherein the active substance with a structure with a structure to formula I-2.

Group V, claim(s) 1-7 (in part) and claims 11-12, drawn to pharmaceutical or diagnostic composition according to claim 1, wherein the active substance with a structure with a structure to formula I-4. If this group is elected, a further election of a single species of active substance with every component named is required.

Group VI, claim(s) 1-7 (in part) and claim 13, drawn to pharmaceutical or diagnostic composition according to claim 1, wherein the active substance with a structure with a

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structure to formula II-2. If this group is elected, a further election of a single species of active substance with every component named is required.

Group VII, claim(s) 1-7 (in part) and claims 14 and 22, drawn to the diagnostic composition according to claim 1, wherein the active substance or at least one of the active substances is labled. If this group is elected, a further election of a single species of active substance with every component named is required.

Group VIII, claim(s) 1 (in part) and claims 17-21, drawn to a method for the treatment or diagnosis of neurodegenerative disorders or amyloid diseases comprising administering a pharmaceutical or a diagnostic composition according to claim 1. If this group is elected, a further election of a single species of active substance with every component named and specific disease (Parkinson syndrome or amlyoid disease) is required.

The groups of inventions listed above do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features because the compounds and methods are distinct from one another because the core structure of each group differs in components, bonding arrangement and chemical properties to such an extent that a reference anticipating any one group among I-VIII would not necessarily render the other groups obvious. The search for each group is not coextensive of the others and separate search and examination must be conducted. Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter, restriction for examination purposes as indicated is proper.:

This application contains claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

Applicant is required, in reply to this action, to <u>elect a single species</u> to which the claims shall be restricted if no generic claim is finally held to be allowable. The reply must also identify the claims readable on the elected species, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered non-responsive unless accompanied by an election.

NOTES: Applicant is respectfully requested to elect a species from with in the elected group above. The elected species should identify, if applicable I-1, I-2, I-3, I-4, I-5, I-6, I-7, I-8, I-9, II-1, II-2, III-1, III-2, III-3, III-4, III-5, III-6, IV-1, IV-2, IV-3, IV-4, IV-5, IV-6, V-1, V-2, V-3, V-4, VI-1, VI-2, wherein R1 to R9 and S1to S3, and other substituents. In addition Applicant is respectfully requested to state which claims are directed to the elected species.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise require all the limitations of an allowed generic claim. Currently, the claim 1 is generic.

REQUIREMENT FOR UNITY OF INVENTION

As provided in 37 CFR 1.475(a), a national stage application shall relate to one invention only or to a group of inventions so linked as to form a single general inventive concept ("requirement of unity of invention"). Where a group of inventions is claimed in a national stage application, the requirement of unity of invention shall be fulfilled only when there is a technical relationship among those inventions involving one or more of the same or corresponding special technical features. The expression "special technical

features" shall mean those technical features that define a contribution which each of the claimed inventions, considered as a whole, makes over the prior art.

The determination whether a group of inventions is so linked as to form a single general inventive concept shall be made without regard to whether the inventions are claimed in separate claims or as alternatives within a single claim. See 37 CFR 1.475(e).

WHEN CLAIMS ARE DIRECTED TO MULTIPLE CATEGORIES OF INVENTIONS

As provided in 37 CFR 1.475(b), a national stage application containing claims to different categories of invention will be considered to have unity of invention if the claims are drawn only to one of the following combinations of categories:

- (1) A product and a process specially adapted for the manufacture of said product; or
 - (2) A product and process of use of said product; or
- (3) A product, a process specially adapted for the manufacture of the said product, and a use of the said product; or
- (4) A process and an apparatus or means specifically designed for carrying out the said process; or
- (5) A product, a process specially adapted for the manufacture of the said product, and an apparatus or means specifically designed for carrying out the said process.

Otherwise, unity of invention might not be present. See 37 CFR 1.475(c).

Due to the complexity of the restriction requirement, a telephone call was not made to request an oral election to the above restriction requirement.

Applicant is advised that the reply to this requirement to be complete must include (i) an election of a species or invention to be examined even though the requirement may be traversed (37 CFR 1.143) and (ii) identification of the claims encompassing the elected invention.

The election of an invention or species may be made with or without traverse. To preserve a right to petition, the election must be made with traverse. If the reply does not distinctly and specifically point out supposed errors in the restriction requirement, the election shall be treated as an election without traverse. Traversal must be presented at the time of election in order to be considered timely. Failure to timely traverse the requirement will result in the loss of right to petition under 37 CFR 1.144. If claims are added after the election, applicant must indicate which of these claims are readable on the elected invention or species.

Should applicant traverse on the ground that the inventions have unity of invention (37 CFR 1.475(a)), applicant must provide reasons in support thereof.

Applicant may submit evidence or identify such evidence now of record showing the inventions to be obvious variants or clearly admit on the record that this is the case.

Where such evidence or admission is provided by applicant, if the examiner finds one of

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the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

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Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to JAGADISHWAR R. SAMALA whose telephone number is (571)272-9927. The examiner can normally be reached on 8.30 A.M to 5.00 P.M.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael G. Hartley can be reached on (571)272-0616. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/J. R. S./ Examiner, Art Unit 1618 /Jake M. Vu/ Primary Examiner, Art Unit 1618

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